Wakefulness Promoting Agents

Nuvigil (armodafinil), Pi	rovigil (mod	dafinil),	Sunosi ((solriamf	etol),	Wakix (pitolis	ant)	
Member	r and Me	dicati	on Info	rmatio	n (req	uired)		
Member ID:		Member Name:						
DOB:			Weight:					
Medication Name/ Strength:			Dose:					
Directions for use:								
	Provider	· Infor	mation	(required)				
Name:	NPI:		Specialty:					
Contact Person:	Office Phone	e:	Offic			e Fax:		
FAX FORM AND RELEVA CHART NOTES a							S,	
Criteria for Approval: Circle the diagnosis ar	nd medication	. (must s	ubmit chart	t notes indic	cating o	one of the following	ng diagnosis)	
Diagnosis, Dose and Age Limitations		Provigi (moda 18 yrs.		Nuvigil (armodafinil) 18 yrs. or older		Sunosi (solriamfetol) 18 yrs. or older	Wakix (pitolisant) 18 yrs. or older	
Daytime somnolence due to obstructive sleep apnea		200mg	/day	150 mg/day		150mg/day		
Narcolepsy		400mg	/day	250mg/day		150mg/day	35.6mg/day	
Narcolepsy with cataplexy							35.6mg/day	
Sedation related to multiple sclerosis treatment		200mg/day						
Shift work sleep disorder		200mg/day		150 mg/day				
Additional criteria for daytime somnolence Patient must use CPAP or prescriber mu Additional criteria for shift work sleep disor Documentation demonstrating excessive Documentation indicating member is we	st submit apporter state of the	ropriate of the control of the contr	clinical ratio		C	nart Note Page #:		
Additional criteria for Sunosi and Wakix: Trial and Failure of modafinil and/or arn Medication:					C	hart Note Page #:		
Dates of therapy:	Details of Failure:							
Re-authorization Criteria: Updated letter of medical necessity or updated letter of the second letter	tive sleep apn ust still be wor	ea, patie rking nigl	nt should co	ontinue on			discontinuation.	
Prescriber's Signature			 Date					

Last updated 4/1/2021